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(54) Preventive or therapeutic agents for Alzheimer's disease, a screening method for Alzheimer's disease, and human tau-protein kinase.

(57) A preventive or therapeutic agent for Alzheimer's disease which comprises a substance exhibiting an inhibitory action to tau-protein kinase I as an effective component is provided. A pharmaceutical composition comprising said agent and a method of inhibiting neuronal cell death in the brain are also provided.

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The present invention relates to a preventive or a therapeutic agent for Alzheimer's disease, a method of screening Alzheimer's disease and tau-protein kinase I which is originated from human being. More particularly, it relates to a preventive or a therapeutic agent for Alzheimer's disease using a tau-protein kinase I inhibitor; a method of screening a preventive or a therapeutic agent for Alzheimer's disease utilizing an amyloid beta-protein; a human-originated tau-protein kinase I which phosphorylates tau-protein, partial peptides thereof or peptides similar thereto; a gene which encodes the kinase; and a method of producing the same.

Alzheimer's disease is a progressive dementia which develops in late middle ages (45 to 65 years old) and its etiological changes are shrinkage of cerebral cortex due to a neuronal cell loss and degeneration of the neurons while, from the pathological view, many senile plaques and neurofibrillary tangles are noted in the brain. There is no pathologically substantial difference between the disease and senile dementia caused by the so-called natural aging which develops in the senile period of 65 years and older ages and, therefore, this is called senile dementia of Alzheimer type.

Numbers of the patients of this disease are increasing with an increase of population of aged people and the disease is becoming serious in the society. There are various theories on the cause of this disease but, the cause has been still ambiguous and, accordingly, there has been a demand for prompt clarification.

It has been known that the quantities which appear in the two pathological changes which are characteristic to Alzheimer's disease and to senile dementia of Alzheimer type are well correlated with the degree of cognitive impairment. Accordingly, studies for clarifying the cause of those diseases by clarifying, in a molecular level, the accumulated insoluble substances resulting in those two pathological changes have been carried out since the first half of the 1980's.

It has been clarified already that a main component of the senile plaques which is one of those pathological changes is amyloid beta-protein (hereinafter, it may be abbreviated as "A $\beta$ P") [Annu. Rev. Neurosci., 12, 463-490 (1989)]. A neurofibrillary tangle which is another pathological change is due to an accumulation of a double-stranded fibrous substance called PHF (paired helical filament) in the neurons and, recently, the components thereof have been identified as ubiquitin and tau-protein which is one of the microtubule-associated proteins characteristic to brain [J. Biochem., 99, 1807-1810 (1986); Proc. Natl. Acad. Sci. USA, 83, 4913-4917 (1986)].

It is believed now that, in Alzheimer's disease, the amyloid beta-protein is extremely accumulated in the neurons and that, as a result of its correlation with the formation of PHF, death of the neurons is resulted.

It has been known that the tau-protein (hereinafter, the protein may be abbreviated as a "tau") is usually a series of related proteins forming several bands at the molecular weights of 48-65 kd on SDS polyacrylamide gel electrophoresis and that it promotes the formation of microtubules.

It has been proved already by the use of polyclonal antibody to PHF [anti-ptau: J. Biolchem. 99, 1807-1810(1986)] and also of monoclonal antibody [tau-1 antibody; Proc. Natl. Acad. Sci. USA, 83, 4913-4917-(1986)] that the tau which is incorporated in the PHF of the brain of Alzheimer's disease is extremely phosphorylated as compared with the normal one.

The present inventors have isolated an enzyme which catalyzes such an abnormal phosphorylation, named it "tau-protein kinase I" (hereinafter, it may be abbreviated as "TPK-I") and clarified its biochemical properties [Seikagaku, vol. 64, no. 5, page 308 (1992)]. The inventors have further cloned the cDNA of rat TPK-I from the cDNA library of cerebral cortex of rats based upon the partial amino acid sequence of TPK-I, whereby the base sequence has been determined and the amino acid sequence has been proposed (Seq. ID No. 2 in the Sequence Listing; Japanese Patent Application 177241/92, FEBS Lett., 325, 167-172 (1993)).

As a result thereof, it has been confirmed that the primary structure of the rat TPK-I is identical with that of the enzyme which is known as a rat GSK-3 $\beta$  (glycogen synthase kinase 3 $\beta$ ) [EMBO J., 9, 2431-2438 (1990)].

However, in finding the drugs which are, effective for the prevention or the therapy of human diseases, the primary structure which are targets for the drug usually vary depending upon the animal species. Therefore, there are many cases that the interaction between the drug and the protein (in other words, sensitivity and effectiveness of the drug) greatly differs depending upon the animal species [e.g. Nature, 360, 161 (1992)]. Thus, in order to find drugs which are really effective to human being, it is desired that the investigation is carried out using proteins which are originated from human being. Particularly in the case of finding the drugs effective for the diseases which have not been found in animals other than human being such as Alzheimer's disease, it is believed to be essential to use proteins originated from human being. However, there has been no report on the separation and purification of TPK-I (or GSK-3 $\beta$ ) from human tissues and, moreover, there has been no report on gene (cDNA) which encodes human TPK-I (or GSK-3 $\beta$ ).

An object of the present invention is to clarify the correlation between the death of neurons and accumulation of PHF and amyloid beta-protein characteristically found in the brains of Alzheimer's disease

and also to apply it to the clarification of the cause of Alzheimer's disease and further to the investigation to the preventive or therapeutic agents therefor.

Another object of the present invention is to clarify the structure of the human-originated TPK-I (which is essential for the progress of investigations for such drugs) on the molecular biological basis and to offer a method of producing it by means of gene technology.

The present inventors have carried out the investigations for achieving the above-given objects and confirmed that, when amyloid beta-protein acted to the neurons in the brain, activity of TPK-I significantly increases whereupon the extremely phosphorylated tau-protein found in PHF of the brains of Alzheimer's disease is resulted and, moreover, the neurons are killed, and that the above-mentioned increase in the TPK-I activity and neuronal cell death in the brain is inhibited by the treatment with the antisense oligonucleotide of the TPK-I.

In addition, with a view that the accumulation of PHF results in the degeneration of neurons in the brains of Alzheimer's disease and successively induces the death, the present inventors have for the first time cloned the gene (which encodes the human-originated TPK-I which is thought to be a key enzyme for the PHF formation) from the cDNA library of human fetus brain, whereupon its primary structure is determined and a method for constant supply (or production) of the human-originated TPK-I has been established.

The present invention has been achieved as a result of the above-given findings followed by further investigations, and its characteristic features are as follows:

- (1) a preventive or a therapeutic agent for Alzheimer's disease, which comprises a substance exhibiting an inhibitory action to tau-protein kinase I as an effective component;
- (2) a preventive or a therapeutic agent for Alzheimer's disease, which comprises an antisense oligonucleotide capable of hybridizing with mRNA or DNA of tau-protein kinase I as an effective component;
- (3) a pharmaceutical composition for prevention or therapy of Alzheimer's disease, which comprises a substance exhibiting an inhibitory action to tau-protein kinase I and a pharmaceutically acceptable carrier;
- (4) a pharmaceutical composition for prevention or therapy of Alzheimer's disease, which comprises an antisense oligonucleotide capable of hybridizing with mRNA and DNA of tau-protein kinase I;
- (5) a method of screening a preventive or a therapeutic agent for Alzheimer's disease in which, when amyloid beta-protein, nerve cells and a drug which is presumed to be effective as a preventive or a therapeutic agent to Alzheimer's disease are incubated and the death of said nerve cells is inhibited, then said drug is judged to be effective as a preventive or a therapeutic agent for Alzheimer's disease;
- (6) a method of inhibiting the death of neurons in the brain, characterized in that a substance which exhibits an inhibitory action to tau-protein kinase I to the neuron in the brain is applied;
- (7) a method of inhibiting the death of neurons in the brain, characterized in that an antisense oligonucleotide which is capable of hybridizing with mRNA or DNA of tau-protein kinase I is applied to the cranial nerve cells;
- (8) human-originated tau-protein kinase I characterized in being represented by an amino acid sequence given in the Seq. ID No. 1 of the attached Sequence Listing or a partial sequence thereof;
- (9) gene which encodes the human-originated tau-protein kinase I which is represented by the amino acid sequence given in the Seq. ID No. 1 of the attached Sequence Listing or a partial sequence thereof;
- (10) recombinant human-originated tau-protein kinase I;
- (11) recombinant vector which is capable of expressing the recombinant human-originated tau-protein kinase I;
- (12) transformant which is obtained by a transformation of the host cells by a recombinant vector which is capable of expressing the recombinant human-originated tau-protein kinase I; and
- (13) a method of producing a recombinant human-originated tau-protein kinase I, characterized in that a transformant obtained by a transformation of host cells by a recombinant vector which is capable of expressing the recombinant human-originated tau-protein kinase I is incubated and then the recombinant human-originated tau-protein kinase I is collected from said culture.

The present invention will be further illustrated as hereunder.

With regard to the substance which exhibits an inhibitory action to tau-protein kinase I in the present invention, any substance will do provided that, when said substance is incubated together with nerve cells and amyloid beta-protein, death of said nerve cells is inhibited. For example, it is chemically-synthesized substance, a substance which is extracted from living cells of microorganisms, etc.

Further, in accordance with the present invention, an antisense oligonucleotide (hereinafter, it may be abbreviated as "TPK-I antisense oligonucleotide") which is capable of hybridizing with mRNA or DNA of

TPK-I is used for prevention or therapy of Alzheimer's disease.

Antisense oligonucleotide is capable of inhibiting the protein synthesis in a level of gene and, therefore, it has been receiving attention in the medical field as a synthetic inhibitor for the proteins causing the disease. The principle is that, when the antisense RNA or the antisense DNA forms a base pair with mRNA in a sense sequence, spread of the gene information is interrupted and synthesis of protein which is the final product is inhibited [Igaku no Ayumi, vol.162, no. 13, 909-911(1992)].

With regard to the TPK-I antisense oligonucleotide applied in the present invention, anything will do provided that it is capable of hybridizing with mRNA or DNA of TPK-I and that it has a sequence for inhibiting the synthesis of TPK-I by, for example, inhibition of transcription, inhibition of splicing of pre-mRNA, inhibition of mRNA septum transmission, inhibition of translation, etc. Usually, that comprising about 15 to 30 nucleotides is used.

Furthermore, the antisense oligonucleotides applicable are a phosphorothioate type in which an oxygen atom which is bonded by means of a double bond with a phosphorus atom at the phosphodiester bond connecting deoxyribonucleosides is substituted with a sulfur atom; a methyl phosphate type in which methyl group is introduced instead of the sulfur atom; a phosphonate type without substitution; and an alpha-oligonucleotide type [Anticancer Drug Des. 6 (66), 606-646 (1991); Anticancer Research, 10, 1169-1182 (1990)]. In addition, in the present invention, it is not always necessary to use a nucleotide type in which a nucleoside derivative is bonded provided that the substance can form a hybrid with the aimed sequence. For example, the antisense compounds which are described in Antisense Research and Development, 1, 65-113 (1991), etc. may be used as well.

Specific examples of the TPK-I antisense oligonucleotides used in the present invention are TPK-I antisense oligonucleotide chain: 5'-TCTCGGTCGCCCGACAT-3' (Seq. ID No. 5 of the Sequence Listing) which is complementary to TPK-I sense oligonucleotide chain: 5'-ATGTCGGGGCGACCGAGA-3' (Seq. ID No. 4 of the Sequence Listing) corresponding to the first six amino acid residues: Met Ser Gly Arg Pro Arg in the translation initiating domain of TPK-I in the primary structure of the rat GSK-3 $\beta$  [same as the primary structure of the rat TPK-I (Seq.ID No. 2 of the Sequence Listing) described in the above-referenced EMBO J., 9, 2431-2438(1990)]; the TPK-I antisense oligonucleotide chain: 5'-TCTGGGCCGCCCTGACAT-3' (Seq. ID No. 7 of the Sequence Listing) which is complementary to the TPK-I sense oligonucleotide chain: 5'-ATGTCAGGGCGGGGCCAGGA-3' (Seq. ID No. 6 of the Sequence Listing) corresponding to the first six amino acid residues: Met Ser Gly Arg Pro Arg in the translation initiating domain of TPK-I in the primary structure of human TPK-I (Seq. ID No. 1 of the Sequence Listing; refer to the examples which will given later); and the like.

The above-mentioned TPK-I sense oligonucleotide and TPK-I antisense oligonucleotide can be easily synthesized by means of commercially-available automatic DNA synthesizers such as a DNA synthesizer manufactured by Applied Biosystems, that manufactured by MilliGen, etc.

As mentioned already, the TPK-I antisense oligonucleotides of the present invention are not particularly limited to those having the above-given sequences provided that they are capable of hybridizing with mRNA or DNA of TPK-I and, so far as the hybrid-forming ability is not deteriorated, a part of the sequence may be substituted with any base. In addition, the antisense oligonucleotides which are changed or modified for passing through a blood-brain barrier as described in Science, 259, 373- 377 (1993) are included in the coverage of the present invention as well.

When the TPK-I antisense oligonucleotides or the substances having an inhibitory action to TPK-I as mentioned above are used as preventive or therapeutic agents for Alzheimer's disease, they may be made into preparations meeting with the particular administering route together with usual carriers. For example, in the case of oral administration, preparations in the form of tablets, capsules, granules, diluted powder, liquid, etc. are prepared.

In preparing solid preparations for oral use, commonly-used fillers, binders and lubricants as well as colorants, disintegrating agents, etc. may be used. Examples of the fillers are lactose, starch, talc, magnesium stearate, crystalline cellulose, methyl cellulose, carboxymethyl cellulose, glycerol, sodium alginate, gum arabic, etc. Examples of the binders are polyvinylalcohol, polyvinyl ether, ethyl cellulose, gum arabic, shellac, white sugar, etc. Examples of the lubricants are magnesium stearate, talc, etc. Besides those, commonly-used ones may be used for the colorants, disintegrating agents, etc. as well. Tablets may be coated by known methods. Liquid preparations may be aqueous or oily suspensions, solutions, syrups, elixirs and the like and may be manufactured by commonly-used methods. In preparing injections, pH-adjusting agents, buffers, stabilizers, isotonic agents, local anesthetics, etc. may be added to TPK-I antisense oligonucleotides or the substances having an inhibitory action to TPK-I and subcutaneous, intramuscular or intravenous injections may be prepared by common methods. With regard to the bases for the manufacture of suppositories, oily ones such as cacao butter, polyethylene glycol, Witepsol (registered

trade mark of Dynamite Nobel) may be used.

Doses of the preparations manufactured as such are not always the same but vary depending upon the symptoms, body weights, ages, etc. of the patients. Usually, however, the amount corresponding to about 1 to 1,000 mg/kg of said drug per day for adults will do and it is preferred to administer by dividing that for 1 to 4 times daily. In some instances, the administration may be carried out once daily to every several or more days.

Examples of the nerve cells used in the present invention are the neuron in the brain collected from mammals and the neuronal cell lines in which the nerve projections are extended by the induction of growth factors such as NGF (nerve growth factor; neurotrophic factor), IGF (insulin-like growth factor), etc. An example of the former is a culture prepared by incubation of tissues of hippocampus of mammals (such as rat) in a complete culture medium. Examples of the latter are PC 12 cells induced by NGF, FGF (fibroblast growth factor), EGF (epidermal growth factor), interleukin 6, etc. [Ann. Rev. Pharma col. Toxicol., 31, 205-228 (1991)]; SH-SY5Y cells induced by IGF [The Journal of Cell Biology, 102, 1949-1954 (1986)]; and those which are disclosed in Cell Culture in the Neurosciences, New York: Plenum Press, pages 95-123 (1955) such as NGF-induced MJB cells, NMB cells, NGP cells, SK-N-SH-SY5Y cells, LAN-1 cells, KA-9 cells, IMR-32 cells and 5-bromodeoxyuridine-induced IMR-32 cells, NMB cells, NGP cells, etc.

Amyloid beta-protein is a main component of senile plaques of Alzheimer's disease and it has been known that said substance is composed of a peptide comprising the following 43 amino acid residues [Science, 250, 279-282 (1990) and Proc. Natl. Acad. Sci. USA, 87, 9020-9023 (1990)].

Amino Acid Sequence of Amyloid beta-Protein (Seq. ID No. 3 of the Sequence Listing):

Asp Ala Glu Phe Arg His Asp Aer Gly Tyr Alu Val His His Gln  
Lys Leu Val Phe Phe Ala Glu Asp Val Gly Ser Asn Lys Gloy Ala  
Ile Ile Gly Leu Met Val Gly Gly Val Val Ile Ala Thr

The present invention will be further illustrated as hereinafter by an example on the behavior of hippocampus cells of rats and the phosphorylation activity of TPK-I when the hippocampus cells were treated with a certain amount of A $\beta$ P and also with TPK-I sense oligonucleotide (hereinafter, referred to as "TPKI-sense") and TPK-I antisense oligonucleotide (hereinafter, referred to as "TPKI-antisense") as controls under certain conditions. When the present invention is carried out as a method of screening the preventive and the therapeutic agent of Alzheimer's disease, hippocampus cells of rat are used as the neurons and, as the agent presumed to be the preventive or the therapeutic agent, TPKI-sense or TPKI-antisense is used.

Certain amount of TPKI-antisense was added to the culture of the hippocampus cells at certain temperature, then certain amount of A $\beta$ P was added thereto, the mixture was kept at certain temperature and the living cell numbers with an elapse of time were measured by a method described in the examples which will be given later. For comparison, the living cell numbers were measured for the case in which only A $\beta$ P was added followed by the same treatments and the case in which TPKI-antisense and A $\beta$ P were added followed by the same treatments. The result showed that, as given in the examples later, the living cell numbers when TPKI-antisense and A $\beta$ P were added were significantly more than those when only A $\beta$ P was added and when TPKI-sense and A $\beta$ P were added and that the TPKI-antisense has an action of inhibiting the death of the cells by A $\beta$ P.

The results of the observations of the samples using a phase contrast microscope (magnifying power: 400) when TPKI-antisense and A $\beta$ P were added to the cell culture followed by allowing to stand for 24 hours, when only A $\beta$ P was added followed by allowing to stand for the same time and when TPKI-sense and A $\beta$ P were added followed by allowing to stand for the same time showed that the cell toxicity by A $\beta$ P was little being similar to the controls only when TPKI-antisense was acted.

Further, the phosphorylation activities of tau-protein by TPKI after 24 hours were measured by the method given in the examples when only A $\beta$ P was added and allowed to stand and when TPKI-antisense and A $\beta$ P were added and allowed to stand same as above. The result was that, as shown in the examples given later, the phosphorylation activity of TPK-I when TPKI-antisense and A $\beta$ P were added was about one half of that when only A $\beta$ P was added and that TPKI-antisense exhibits an activity of inhibiting the phosphorylation activity of TPK-I.

Out of the above results, it may be concluded that, when the present invention is carried out as a method of screening the preventive and the therapeutic agent for Alzheimer's disease, TPKI-antisense is

effective as said preventive and therapeutic agent. Incidentally, the effectiveness of the agents other than the TPK-I antisense oligonucleotide can be evaluated similarly.

Now, the method of obtaining the human-originated TPK-I and the method of production thereof will be illustrated as hereunder.

5 The TPK-I originated from human being of the present invention may, for example, be manufactured as follows. Thus, microtubule fractions were obtained from an extract of human brain immediately after death by means of temperature-depending polymerization and depolymerization and then, operations such as phosphocellulose column chromatography, gel filtration, hydroxyapatite column chromatography, S-Sepharose column chromatography, heparin column chromatography, etc. are combined according to a  
10 method by Uchida, et al. [Seikagaku, vol.64, no.5, page 308 (1992)] whereby pure protein is obtained. The (partial) primary structure of such a pure protein may be determined by conducting a conventional amino acid analysis. It is not easy to obtain the human brain tissues in large quantities and it is difficult to purify the human TPK-I and, therefore, it is also possible that, by a method which will be given later, gene is previously cloned and the amino acid sequence is deduced therefrom whereby the primary structure is  
15 determined.

The human TPK-I of the present invention prepared as such is a protein in which the primary structure is represented by the amino acids described in the Seq. ID No. 1 of the Sequence Listing (420 amino acid residues; molecular weight: 46,719; isoelectric point: 9.21) and alterations such as removal, substitution, modification or addition of some amino acids may be carried out within such a range that the functions  
20 (action, substrate specificity, etc.) will not be deteriorated.

The gene (cDNA) which encodes the above TPK-I may be cloned by such a method that in which the corresponding protein is purified from natural material, its partial amino acid sequence is determined and the DNA probe corresponding thereto is utilized; that in which homology with the protein of the same species or the corresponding protein of the different animal species is utilized; that in which an antibody  
25 which is specific to the corresponding protein is utilized; that in which a detection of the specific function or the protein is utilized; etc. The present inventors have previously purified TPK-I from an extract of brain of rat or bovine and, depending upon the information of the partial amino acid sequence thereof, they cloned the rat TPK-IcDNA from the rat brain cDNA library (Seq. ID No. 2 of the Sequence Listing; Japanese Patent Application No. 177241/92, FEBS Lett., 325, 167-172 (1993)).

30 Usually, however, the homology of the primary structures of rats with human beings in the same protein is, in most cases, around 90% or more and, therefore, it is possible to clone human TPK-IcDNA from rat TPK-IcDNA by utilizing said homology. Thus, lambda-phage is infected to Escherichia coli by a method of Tomizawa, et al. ["Experiments in Bacteriophage" (Iwanami Shoten), pages 99-174 (1970)] from cDNA library containing the gene which encodes human TPK-I such as human fetus brain cDNA library followed  
35 by culturing. The plaques formed thereby were selected by a plaque hybridization method ["Molecular Cloning" Cold Spring Harbor Laboratory, pages 320-328 (1982)] using a rat TPK-IcDNA or DNA fragments having a partial structure thereof as a probe. The phage is promulgated from positive plaques by a method of Tomizawa, et al., then DNA is prepared by a method of T. Maniatis, et al. ["Molecular Cloning", Cold Spring Harbor Laboratory, page 85 (1982)] or after subjecting to a subcloning if necessary, cleaved by a  
40 suitable restriction enzyme such as EcoRI and cloned to a plasmid such as pUC18 or pUC19. As such, cDNA of human TPK-I is prepared and its base sequence can be determined, for example, by a dideoxy method of Sanger, et al. [Proc. Natl. Acad. Sci. USA, 74, 5463 (1977)].

An example of the above-mentioned gene (cDNA) encoding the human-originated TPK-I is that which has a base sequence given in the Seq. ID No. 1 of the sequence Listing.

45 In the human TPK-I prepared by the present invention, 5 amino acid residues were different out of the 420 amino acid residues in the amino acid sequence of the rat TPK-I and the homology in the bases in the translation domain was 92.5%.

The above-prepared human TPK-I or its partial peptides can be expressed and generated by means of gene recombination. Thus, the 5'-terminal of the human TPK-IcDNA or its fragment is modified or added  
50 with precursor gene, inserted into the downstream of the promoter of the expressed vector and then the vector is induced into host cells such as bacteria, yeasts, insect cells, animal cells, etc. The transformed host cells as such is cultured under a suitable condition whereby the recombinant human TPK-I is produced in or outside the cells.

55 Examples of the transformable host cells are bacteria (procaryotic cells) such as Escherichia coli (K-12 strain), Bacillus subtilis, etc.; yeasts such as Saccharomyces cerevisiae; insect cells such as ovary-originated cells (Sf9 cell strain) of Spodoptera spp.; and (mammalian) animal cells such as ovary-originated cells (CHO cells) of Chinese hamster, mouse C127 cells, kidney-originated cells (COS cells) of African green monkey, mouse L cells, mouse FM3A cells, kidney-originated cells (HEK cells, 293 cells) of human

fetus, etc.

The expression vector which is suitably used is that which contains a promoter at the position where a gene (cDNA) coding the human TPK-I or the DNA fragment thereof can be transcribed. For example, when cells are used as a host, it is preferred that the expression vector is composed of promoter, ribosome binding (SD) sequence, human TPK-I-encoding gene or fragment thereof, transcription terminating factor and promoter-controlling gene. Even when eucaryotic cells such as (mammalian) animal cells, insect cells, yeasts, etc. are used as host cells, the fundamental unit comprising the expression vector is the same as that in the case of the above-mentioned bacteria.

Examples of the promoter when bacteria are used as host cells are those originated from *Escherichia coli*, phage, etc. such as lactose operon (lac), tryptophan-synthesizing enzyme (trp), lambda-phage P<sub>L</sub>, *E. coli* T7 polymerase promoter and tac (hybrid promoter originated from trp and lac UV5). The examples in the case of yeasts are promoters of gene of enzymes such as phosphoglyceric acid kinase (PGK), glyceraldehyde-3-phosphoric acid dehydrogenase (GPD), repressible acidic phosphatase (PHO5) and alcohol dehydrogenase 1 (ADH1). The examples in the case of insect cells are used as the host are promoter of polyhedron gene of baculovirus, etc. The examples in the case of (mammalian) animal cells are SV40 initial promoter, SV40 late promoter, apolipoprotein E gene promoter, etc.

Examples of the ribosome binding sequence are those which are originated from *E. coli*, phage, etc. and those which are partially complementary to the base sequence of the 3'-terminal domain of 16S ribosome RNA.

Though the transcription terminating factor is not always necessary, it is preferred to have that which is rho<sub>p</sub>-independent such as lipoprotein terminator, trp operon terminator, etc.

With regard to the sequence of those factors necessary for the expression on the expression plasmid (vector), it is desired that they are placed in the order of promoter, SD sequence, human TPK-I-encoding gene or fragment thereof and transcription terminating factor from the 5'-upstream side.

Specific examples of the expression vectors satisfying those requirements in case that bacteria are used as the host are pKK233-2 (manufactured by Pharmacia) and pET3C [Gene, 56, 125 (1987)]. Expression vector pGEX series (Pharmacia) which are expressed as fused protein may be used as well in the same manner. When yeasts are used as a host, the vector in which the above-mentioned promoter (and, further, gene which complements the auxotrophic mutant as a selected marker such as trp1 and leu2) is incorporated into YEp vector having a replication origin of 2 micron DNA is suitably used. In the case of insect cells, an example is Maxbac (trademark) which is a baculovirus expression system manual version 1.4 of Invitrogen. In the case of animal cells, those having the above-mentioned promoter and selective marker gene such as neomycin-resisting gene (Neo) and dihydrofolic acid reductase gene (DHFR) are suitably used. When eucaryotic cells are used as a host, a shuttle vector to *E. coli* may be used as well.

Transformation of host cells may be carried out by conventional manner.

Culture of the transformant may be carried out by a method depending upon a method of T. Maniatis, et al. described in "Molecular Cloning" Cold Spring Harbor Laboratory, 1982. Though the culture temperature is not always the same depending upon the conditions such as the host cells, a temperature of about 25 to 40 °C is suitable.

The human TPK-I produced by such host-vector systems can be purified by a series of purifying steps corresponding to hosts and culturing conditions such as extraction from the host, salting-out and chromatography using various columns. In the case of column chromatography, the suitably-used ones are phosphocellulose column chromatography, hydroxyapatite column chromatography, S-Sepharose column chromatography, heparin column chromatography, blue Sepharose column chromatography, etc.

The recombinant human TPK-I prepared as such is capable of phosphorylating proteins such as tau-protein, glycogen-synthesizing enzyme, protooncogene product c-jun, etc. and partial peptides thereof and the progress of the phosphorylation can be confirmed by, for example, the conditions as given below. Thus, the recombinant human TPK-I of the present invention is added, together with a suitable amount of the substrate protein, to a buffer of pH. 5.0-8.0 containing 0.2-4.0mM magnesium acetate and 0.2-4.0mM adenosine triphosphate, the mixture is incubated at the room temperature to 40 °C and the phosphorylation of the substrate protein is checked and determined by radiochemical, proteinchemical or immunochemical means. Consequently, then an agent is added to this reaction system and the resulting promotion or inhibition of the phosphorylation reaction is checked, it is possible to find the agent having a physiologically important meaning whereby the investigation on the agent which is effective for the prevention or the therapy of human disease is now possible.

## Examples

The present invention will be illustrated by way of the following examples though the present invention is not limited to those examples so far as they are not out of the characteristic feature of the present invention. Incidentally, judgement of the cytotoxicity, measurement of the phosphorylation of tau-protein and immunohistochemistry by Alz-50 antibody were carried out in accordance with the following methods. Further, in each of the following examples, at least three independent experiments were carried out and the data were given by their average values.

## Judgement of Cytotoxicity:

Numbers of many normal and healthy cells were counted by a phase contrast microscope as an index of the living cells after the treatment. Normal cells mean those which have morphologically flat circumference and many nerve cell projections while the degenerated cells were judged by checking the irregular shape, degeneration of the neural projections, etc. Numbers of the living cells here counted in a well. In the standard culture liquid, the cell numbers were not less than 400 per well. The result was confirmed by an immunohistochemical means.

## Measurement of Degree of Phosphorylation of tau-Protein:

Hippocampus cells were collected from the culture medium by washing with an ice-cooled phosphate buffer for three times. The cells were suspended in a buffer A (pH: 6.8) which contained 1mM EGTA, 0.5mM magnesium acetate and 20mM 2-(N-morpholino)-ethanesulfonic acid containing a phosphatase inhibitor (1mM okadaic acid; manufactured by Seikagaku Kogyo) and a protease inhibitor (1mM phenylmethylsulfonyl fluoride and each 1 micro gram/ml of leupeptin, pepstatin and aprotinin), homogenized and centrifuged at 14,000 rpm for one hour and the supernatant liquid was used for checking the phosphorylation.

The rat tau-protein expressed in E. coli BL21 by a gene recombination was purified by a method described in J. Biol. Chem., 267, 10897-10901 (1992).

The hippocampus extract (1 microliter) was added to a solution of the rat tau-protein (400 micrograms/ml) dissolved in a buffer A containing 1mM [ $\gamma$ - $^{32}$ P]ATP (10-20 Ci/mmol) and then 10 micromoles of okadaic acid was added to make the final volume 10 microliters. This was incubated at 37°C for three hours and the reaction was stopped by adding a buffer for electrophoresis. After subjecting to a 10% polyacrylamide gel electrophoresis, the  $^{32}$ P in the tau-protein was observed by a laser image analyzer (Fuji BAS 2000).

## Immunohistochemistry by an Alz-50 Antibody:

The cultured medium of the hippocampus cells was fixed in a phosphate buffer for ten minutes using 4% paraformaldehyde. The fixed culture liquid was incubated for 30 minutes in a Tris buffer containing 0.2% Triton X-100 so that the cells were made permeable.

Then this culture medium was subjected to an immunolabelling using a 1:5 diluted Alz-50 mouse monoclonal antibody [Science, 232, 648-650(1986)], Vectastain ABC avidin-biotin-enzyme peroxide detector kit (manufactured by Vector Laboratory) and diaminobenzidine tetrahydrochloride as a dye.

## Example 1.

## Preparation of Culture Medium of Cells:

The primary culture medium of hippocampus of rats was prepared in accordance with a method described in Brain Res., 126, 397-425 (1977). Thus, the hippocampus tissues were collected from embryo of the rats of 18 days after fertilization and digested in papain (protease) (10 U/ml) at 37°C for 20 minutes. The resulting cells were added to a Dulbecco's modified Eagle's medium supplied with 5% bovine fetus serum, 5% horse serum, 10 micrograms/ml insulin, 0.1 mg/ml transferrin, 1 microgram/ml aprotinin, 1 mM sodium pyruvate and 84 micrograms/ml gentamycin. This was planted to a well for tissue culture covered with poly-L-lysine at the density of  $2 \times 10^5$  cells/cm<sup>2</sup>, cultured for three days and treated with 1 micromole of cytosine-beta-arabinofuranoside for 24 hours and the cells of the fifth day of the culture were used.



Preparation of A $\beta$ P:

A $\beta$ P peptide (Seq. ID No. 3 of the Sequence Listing) comprising the already-mentioned 43 amino acid residues was synthesized by a method which was described in Science, 250, 279-282(1990) and Proc. Natl. Acad. Sci. USA, 87, 9020-9023(1990) and, after being purified, it was dissolved in 35% acetonitrile to prepare a stock solution of 2M.

## Preparation of TPKI-Sense and TPKI-Antisense:

Rat GSK-3 $\beta$  [EMBO J., 9, 2431-2438(1990)], i.e. the TPKI-sense comprising the following 18 bases corresponding to the translation initiating domain of the primary structure of rat TPK-I (FEBS Lett., 325, 167-172 (1993)) and the TPKI-antisense which is complementary thereto were synthesized using an automatic DNA synthesizer (MilliGen), recovered from 20% acrylamide-urea gel and purified by means of an ethanol precipitating method and the precipitate was dissolved in water to adjust to a concentration of 1 micromole.

TPKI-Sense: 5'-ATGTCGGGGCGACCGAGA-3' (Seq. ID No. 4 of the Sequence Listing)

TPKI-Antisense: 5'-TCTCGGTCGCCCCGACAT-3' (Seq. ID No. 5 of the Sequence Listing)

## Inhibiting Action for the Death of Cranial Nerve Cells:

The culture medium of the hippocampus prepared by the above-mentioned method was subjected to the following treatments (b) to (d), numbers of the living cells with an elapse of time were counted and the result is given in Table 1.

(a) Nontreated culture medium (control):

(b) TPKI-antisense (1 micromole) was added to 1 ml of the cell culture medium and, after five minutes, 20 micromoles of A $\beta$ P was added followed by keeping at 37°C for 24 hours.

(c) A $\beta$ P (20 micromoles) was added to 1 ml of the cell culture medium followed by keeping at 37°C for 24 hours.

(d) TPKI-sense (1 micromole) was added to 1 ml of the cell culture medium and, after five minutes, 20 micromoles of A $\beta$ P was added followed by keeping at 37°C for 24 hours.

Table 1

Treating Agents	Numbers of Living Cells (%) After	
	6 hours	21 hours
(Control)	100	100
A $\beta$ P + TPKI-Antisense	83.0	72.6
A $\beta$ P	41.3	25.4
A $\beta$ P + TPKI-Sense	49.5	17.1

Table 1 shows the numbers of the living cells with an elapse of time after the above-mentioned treatments (b), (c) and (d) and the numbers are given in terms of percentages to the control.

As shown in Table 1, the numbers of the living cells after 6 and 21 hours of the treatment of the hippocampus cells with TPKI-antisense and A $\beta$ P (b) were significantly more than those of the case treated only with A $\beta$ P (c) and of the case treated with TPKI-sense and A $\beta$ P (d). This fact clearly shows that the TPKI-antisense significantly inhibits the death of the cells by A $\beta$ P.

Further, it was clarified by the observations of the above-mentioned cases of (b) to (d) after 24 hours using a phase contrast microscope (magnifying power: 400) that, only in the case of (b) where TPKI-antisense and A $\beta$ P were acted to the hippocampus cells, the cytotoxicity by A $\beta$ P was little and similar to the case of the control.

## Phosphorylation of tau-Protein:

Phosphorylating activity of the TPK-I was measured by the above-mentioned method for the samples of (1) untreated cell culture medium (control); (2) a sample in which 1 micromole of TPKI-antisense was added to 1 ml of the cell culture medium followed by adding 20 micromoles of A $\beta$ P after 5 hours; and (3) a sample in which 20 micromoles of A $\beta$ P was added to 1 ml of the cell culture medium and the result is

given in Table 2. The phosphorylating activity of TPK-I in Table 2 shows that (units/mg protein) per mg of the protein in the supernatant liquid wherein one unit is equivalent to the intensity of the radioactivity measured by a laser image analyzer (BAS 2000; Fuji).

Table 2

Treating Agent	Phosphorylating Activity of TPKI (unit/mg protein)
(Control)	39.6
A $\beta$ P + TPKI-Antisense	31.6
A $\beta$ P	66.2

As shown in Table 2, the phosphorylating activity of the case (2) in which TPKI-antisense and A $\beta$ P were acted on the cell culture medium was only about one-half of that of the case (2) in which only A $\beta$ P was acted. Thus, it is clear that the TPKI-antisense significantly inhibits the phosphorylating activity of TPK-I by A $\beta$ P.

#### Example 2. Cloning of Human TPK-IcDNA.

Commercially-available human fetus brain cDNA library (prepared by inserting a 1:1 mixture of cDNA synthesized from mRNA of human fetus brain using oligo dT and random primer to lamda-ZAPII; manufactured by Strategen) was infected to a host which was E. coli XL1-blue [W. O. Bullock, et al: Biotechnology, 5, 376-379(1987)] to form plaques. The plaques (numbers: 450,000) were screened using a probe which was prepared by a part of the translation domain (170 base pair from the 1137th HindIII site to the 1306th A; Seq. ID No. 8 of the Sequence Listing) of the rat TPK-IcDNA (Seq. ID No. 2 of the Sequence Listing) whereby 19 positive clones were obtained. Among those, two clones were subcloned to Bluescript SK (Strategen) and then EcoRI fragments which hybridize with the above probe were subcloned to vector pUC19 [C. Yanisch-Perrou, et al: Gene, 33, 103 (1985)]. The restriction enzyme map of the EcoRI fragments of the two clones prepared as such is given in Fig. 1.

Base sequence was determined for entire domains of the clone #1 by a dideoxy method and it lacked the N-terminal moiety of human TPK-I protein. Total length of clone #2 was 2.2 kilobases and, out of a comparison with the restriction enzyme map, it was presumed to probably contain clone #1. Therefore, the base sequences corresponding to 5'-untranslated domain and N-terminal domain mostly comprising human TPK-I protein were determined by a dideoxy method of Sanger, et al. The base sequence of cDNA out of the both results and the amino acid sequence of TPK-I supposed therefrom are given in Seq. ID No. 1 of the Sequence Listing.

Further, comparison with the amino acid sequence of the rat TPK-I is given in Fig. 2.

#### Example 3. Expression of Human TPK-I by Insect Cells.

NruI-EcoRI fragments containing entire length of translation domain of human TPK-IcDNA were inserted to a SmaI-EcoRI part of transfer vector PVL1392 [Invitrogen; N. R. Webb and M. D. Summers: Technique, 173-188(1990)] prepared by insertion of virus-originated DNA fragments containing baculovirus (nuclear polyhedrosis virus) polyhedron gene and promoter thereof into vector PUC8 (E. coli-hosted plasmid vector) whereupon an expression vector PVL-TPKI was prepared.

Cell strain Sf9 originated from ovalium cells of Spodoptera spp. was cultured in a medium for insect cells FNM-FH [This was prepared as follows; thus, 0.35mg/lit of sodium bicarbonate (manufactured by Wako Pure Chemical), 3.3 mg/ml of TC lactalbumin hydrolysate (manufactured by Difco) and 3.3 mg/ml of TC yeast late were added to a Grace's insect medium (Sigma), adjusted to pH 6.2, sterilized and then a heat-processed 10% bovine fetus serum, 50 micrograms/ml of gentamycin sulfate and 2.5 micrograms/ml of amphotericin B were added thereto.] and co-infected with wild baculovirus DNA and vector DNA whereupon a homogeneous recombination between them took place to some extent and, as a result, recombinant virus having a TPK-I expressing system was prepared.

Selection of the wild virus-infected cells and the recombinant virus-infected cells was conducted visually and, as a result of repeated selections for three times, the recombinant virus-infected cells were separated. The virus-containing liquid with high infectivity obtained from the supernatant of the cells was further infected to Sf9 cells and cultured for 72 hours to recover  $5 \times 10^8$  cells. They were suspended in 30 ml of a

buffer A [comprising 10 mM sodium phosphate (pH: 7.05), 1 mM ethylenediaminetetraacetic acid, 5mM ethyleneglycol bis (2-amino-ethyl ether) tetraacetate, 2mM dithiothreitol, 10 mM magnesium chloride, 0.1mM sodium orthovanadate, 40 micrograms/ml phenyl methanesulfonyl fluoride, 1 microgram/ml leupeptin, 1 microgram/ml pepstatin and 1 microgram/ml antipain], homogenized and centrifuged at 105 G to  
 5 recover the supernatant liquid. Then the supernatant was subjected to a phosphocellulose column chromatography (filled with P-11; Whatman) and fractionated with a buffer B [comprising 25mM tris-(hydroxymethyl)aminomethane hydrochloride (pH: 7.5), 1mM ethylenediaminetetraacetic acid, 1mM dithiothreitol, 0.1% beta-mercaptoethanol, 5% glycerol and 50 mM sodium chloride] with a gradient of sodium chloride concentrations of 50mM to 250 mM. Each fraction of the eluate was analyzed by means of an  
 10 electrophoresis and an immunoblot technique and an anti-TPK-I antibody positive fraction which cross-reacts with an anti-rat TPK-I amino terminal antibody [rabbit antiserum obtained as an antigen by expressing rat TPK-I amino terminal 36 residues (Seq. ID No. 9 of the Sequence Listing) as an E. coli-hosted vector system as its fused protein with beta-galactosidase] was pooled. This was concentrated by means of an ultrafiltration, subjected to a blue sepharose column chromatography (filled with Blue-Sephadex CL-6B of Pharmacia) and fractionated by a buffer C [comprising 20 mM N-2-hydroxyethyl-piperazine-N'-2-ethanesulfonic acid (pH: 7.5), 1 mM ethylenediaminetetraacetic acid, 0.1 mM dithiothreitol and 5% glycerol] with a sodium chloride concentration gradient of 0 to 1M. Anti-TPK-I antibody positive  
 15 fraction was pooled by analysis of immunoblotting and electrophoresis for each fraction, subjected to an ultrafiltration and dialyzed against a buffer D [comprising 100 mM 2-(N-morpholino)-ethanesulfonic acid (pH: 6.5), 0.5 mM magnesium acetate, 1 mM ethyleneglycol bis(2-aminoethyl ether) tetraacetate, 10% glycerol, 0.02% polyoxyethylenesorbitan monolaurate (Tween 20), 0.1 mM phenylmethanesulfonyl fluoride, 1 microgram/ml pepstatin, 1 microgram/ml antipain, 1 microgram/ml leupeptin and 5 mM beta-mercaptoethanol] to give 1 ml of enzyme liquid. Total protein obtained was 0.4 mg.

Progress of phosphorylation was checked using this enzyme solution by the following two methods.

25

#### (Phosphorylating Method I)

tau-Protein extracted from bovine brain followed by purification (2 microliters; 1.5 mg/ml concentration) and 1 microliters of the above partially-purified enzyme solution were mixed. To the mixture was added a  
 30 solution containing 2mM adenosinetriphosphate and 2mM magnesium acetate and [gamma-32P]-adenosinetriphosphate so that the phosphorylation of tau-protein was conducted at room temperature for 20 hours whereby the amount of phosphoric acid incorporated in tau-protein was evaluated.

#### (Phosphorylating Method II)

35

Phosphorylation reaction which was the same as in the method I was conducted with an exception that no [gamma-32P]adenosinetriphosphate was contained followed by subjecting to an SDS electrophoresis to blot to nitrocellulose. The blotted tau-protein was subjected to an immunodyeing with anti-tau antibody (rabbit antiserum to chicken fetus brain-originated tau-protein) and anti-p-tau antibody [Ihara, et al: J. Biochem., 99, 1807-1910(1986)].  
 40

As a result of the method I, incorporation of tau-protein into phosphoric acid was confirmed while the result of the method II was that:

- 1) mobility of tau-protein after the reaction was less than that of tau-protein which was not phosphorylated; and
- 2) tau-protein which was not phosphorylated did not react with anti-p-tau antibody while tau-protein after the reaction reacted with anti-p-tau antibody.

45

Those results indicate that the outcome was the same as that in the phosphorylation of tau-protein using the TPK-I purified from animal brain.

#### 50 Example 4. Phosphorylation of Peptide by Recombinant Human TPK-I.

Peptide (hereinafter, abbreviated as "K2") represented by the amino acid sequence described in the Seq. ID No. 10 of the Sequence Listing was synthesized. This peptide was phosphorylated by the same manner as in the phosphorylating method II in Example 3 with an exception that tau-protein kinase II (TPK-II) purified from bovine brain microtubule was used instead of TPK-I whereupon phosphorylated peptide (hereinafter, abbreviated as "p-K2") was obtained.  
 55

Phosphorylations of K2 and p-K2 were conducted according to a phosphorylating method I of Example 3 using human-originated partially purified TPK-I prepared in Example 3 whereupon the progress of

phosphorylation of p-K2 was clearly noted while phosphorylation of K2 was slow and its initial speed was about one-tenth of that of p-K2.

The result shows that the outcome was the same as the phosphorylation of K2 and p-K2 using TPK-I purified from animal brain.

5

#### Example 5. Expression of Recombination of Human TPK-I by Escherichia coli.

SacI-EcoRI fragment of human TPK-IcDNA clone #2 obtained in Example 2 was introduced into an SacI-EcoRI part of vector PUC19 [C. Yanisch-Perrou, et al: Gene, 33, 103 (1985)] to prepare pUSE2. In the  
10 meanwhile, in order to prepare an NdeI part in an oligonucleotide from 598th to 629th members of cDNA represented by the base sequence described in the Seq. ID No. 1 of the Sequence Listing, a plus strand oligonucleotide (Seq. ID No. 11 of the Sequence Listing) wherein CAT was inserted between 613th and 615th member and a minus strand oligonucleotide (Seq. ID No. 12 of the Sequence Listing) from 1076th to 1047th members were synthesized and a cDNA fragment (Seq. ID No. 13 of the Sequence Listing) ranging  
15 from 598th to 1076th members and having NdeI part duplicating with the initiation codon was obtained by a PCR method [Saiki, et al: Nature, 324, 126 (1986)].

Fragments ranging from a 5'-terminal of the cDNA fragment obtained by a PCR method to a SacI part were inserted to SmaI-SacI part of pUSE2. The NdeI-EcoRI fragment of the plasmid vector was introduced into NdeI-BamHI part of pET3C [A. H. Rosenberg, et al: Gene, 56, 125(1987)] which is one of the vectors  
20 having E. coli T7 polymerase promoter to construct pET3C/TPKI.

pET3C/TPKI was transformed by conventional means using E. coli BL21 (DE3) [F. W. Studier and B. A. Moffatt: J. Mol. Biol. 189, 113 (1986)] as a host to prepare a recombinant. The resulting recombinant E. coli was cultured at 37°C until the middle stage of the logarithmic growth phase, kept at 21°C, 0.3mM (final concentration) of IPTG (isopropyl-beta-D(-)-thiogalactopyranoside) was added and cultured for four hours  
25 more. The living cells (5 g) were suspended in 50 ml of a buffer E [comprising 20 mM of 2-(N-morpholino)-ethanesulfonic acid (pH: 6.5), 1 mM of ethylenediaminetetraacetic acid, 5mM of beta-mercaptoethanol and 50 mM of sodium chloride], disintegrated with ultrasonic wave and centrifuged at 100,000 g for one hour. The supernatant was subjected to a phosphocellulose column chromatography (filled with P-11; Whatman) which was balanced with a buffer E and then subjected to a gradient elution with sodium chloride concentrations of 50 to 500 mM whereby the fraction which was positive to anti-TPK-I antibody was pooled and concentrated. This was dialyzed against a buffer F [comprising 20 mM of N-2-hydroxyethyl-piperazine-N'-2-ethanesulfonic acid (pH: 7.2), 1 mM of ethylene-diaminetetraacetic acid and 5 mM of beta-mercaptoethanol], subjected to a blue sepharose column chromatography (filled with Blue-Sepharose CL-6B of Pharmacia) and eluted with a concentration gradient of 0 to 1M of sodium chloride. Anti-TPK-I antibody  
30 positive fractions were collected and dialyzed against a buffer D.

Phosphorylation of tau-protein was conducted by the phosphorylating methods I and II by the same manner as in Example 3 using the resulting partially-purified TPK-I. It was found that, as a result of the phosphorylating method I, 1.2 moles of phosphoric acid was incorporated into one molecule of tau-protein while, as a result of the phosphorylating method II, the mobility of electrophoresis of tau-protein after the  
40 reaction became small and the reaction with anti-ptau antibody became positive.

When the partially-purified TPK-I was used for phosphorylation of the peptides K2 and p-K2 by the same manner as in Example 4, the phosphorylation of p-Ka proceeded while that of K2 hardly proceeded.

Those results show that the recombinant TPK-I prepared in this example had the same property as that of TPK-I purified from animal brain and of recombinant TPK-I prepared in Example 3.

45

#### (Merit of the Invention)

In accordance with the preventive and the therapeutic agent of Alzheimer's disease of the present invention, the phosphorylating activation of tau-protein kinase I by amyloid beta-protein was inhibited  
50 whereby the death of the neuron in the brain can be inhibited. Further, it is possible to conduct a screening of the preventive or the therapeutic agent of Alzheimer's disease utilizing the above mechanism.

Moreover, the human-originated TPK-I of the present invention is an enzyme which specifically acts to tau-protein which is suggested to be related to Alzheimer's disease and also to senile dementia of Alzheimer's disease type and, therefore, its application to clarification of cause of those diseases and to the  
55 investigations for the agents for the prevention and the therapy thereof can be expected.

Brief Explanation of the Drawings:

Fig. 1 is a drawing which shows the restriction enzyme map of the human TPK-I.

Fig. 2 is a drawing which shows the comparison of amino acid sequences of human TPK-I and rat TPK-

5 I. In the drawing, each amino acid is represented by a single letter.

SEQUENCE LISTING

10 (1) GENERAL INFORMATION:

(i) APPLICANT: MITSUBISHI KASEI CORPORATION

(ii) TITLE OF INVENTION: PREVENTIVE OR THERAPEUTIC  
15 AGENTS FOR ALZHEIMER'S DISEASE, A SCREENING METHOD FOR  
ALZHEIMER'S DISEASE, AND HUMAN tau-PROTEIN KINASE

(iii) NUMBER OF SEQUENCES: 13

20 (iv) CORRESPONDENCE ADDRESS:

(A) ADDRESSEE:

(B) STREET: 5-2, Marunouchi 2-chome, Chiyoda-ku

(C) CITY: Tokyo

(E) COUNTRY: Japan

25 (F) ZIP: 100

(v) COMPUTER READABLE FORM:

(A) MEDIUM TYPE: Floppy disk

(B) COMPUTER: IBM PC compatible

30 (C) OPERATING SYSTEM: PC-DOS/MS-DOS

(D) SOFTWARE: PatentIn Release #1.0, Version #1.25

(vi) CURRENT APPLICATION DATA:

(A) APPLICATION NUMBER:

35 (B) FILING DATE:

(C) CLASSIFICATION:

(viii) ATTORNEY/AGENT INFORMATION:

(A) NAME:

40

(2) INFORMATION FOR SEQ ID NO:1:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2088

45 (B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA to genomic RNA

50

(vi) ORIGINAL SOURCE:

(A) ORGANISM: human being

55

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:1:

5 TTACAGGTGT GAGCCACCTC GCCCAGCTGA GTTCAGTATA ATTTTCAATG AGAAACTGAA 60  
 ATTCAGTTTT ATAATCAAAG AGCATGTTTG CTGAAGCCAT CATTCTCAGC AAATAATAC 120  
 AGGGACAGAA AACCAAACAC CGCATGTTCC ACTCATAAGT GGGAGTTGAA CAATGAGAAC 180  
 10 ACACGGACAC AGGGAGGGAA ACATCACACA CCAGGGCCTG TCAGGCGGTC AGGGGTAAGG 240  
 GGAGAGAGAG CATCGAGACA AATATCTAAG GTATGCGGGG CTAAAACCT AGATGATGGT 300  
 TGATAGGTGC AGCAAACCAC CATGGCACAT GTATACCTGT GTAACAAACC CGCACGTCCT 360  
 15 GCACATGCAT CCCACAACCT AAAGCAAAAT AAAAATATAT ATATTTTCA TATTTTCATA 420  
 TATAATATAT AAATATATAA TTAAGATAAA ATATTACATA TTACATATGT ATAAATTCAT 480  
 20 ATATAACATA TAAAATATAT AATATTATAT ATTATATACA TGTGTATATA AAATCTGGCT 540  
 GCGGAGTTTT TGATCTATAC ATTGAACAAA TTGTCTCACC TACTGATGAA AAGGTGATTC 600  
 GCGAAGAGAG TGATC ATG TCA GGG CGG CCC AGA ACC ACC TCC TTT GCG GAG 651  
 25 Met Ser Gly Arg Pro Arg Thr Thr Ser Phe Ala Glu  
 1 5 10  
 30 AGC TGC AAG CCG GTG CAG CAG CCT TCA GCT TTT GGC AGC ATG AAA GTT 699  
 Ser Cys Lys Pro Val Gln Gln Pro Ser Ala Phe Gly Ser Met Lys Val  
 15 20 25  
 35 AGC AGA GAC AAG GAC GGC AGC AAG GTG ACA ACA GTG GTG GCA ACT CCT 747  
 Ser Arg Asp Lys Asp Gly Ser Lys Val Thr Thr Val Val Ala Thr Pro  
 30 35 40  
 40 GGG CAG GGT CCA GAC AGG CCA CAA GAA GTC AGC TAT ACA GAC ACT AAA 795  
 Gly Gln Gly Pro Asp Arg Pro Gln Glu Val Ser Tyr Thr Asp Thr Lys  
 45 45 50 55 60  
 CTC ATT GGA AAT GGA TCA TTT GGT GTG GTA TAT CAA GCC AAA CTT TGT 843  
 Leu Ile Gly Asn Gly Ser Phe Gly Val Val Tyr Gln Ala Lys Leu Cys  
 50 65 70 75  
 55

EP 0 616 032 A2

	GAT TCA GGA GAA CTG GTC GCC ATC AAG AAA GTA TTG CAG GAC AAG AGA	891
5	Asp Ser Gly Glu Leu Val Ala Ile Lys Lys Val Leu Gln Asp Lys Arg	
	80 85 90	
	TTT AAG AAT CGA GAG CTC CAG ATC ATG AGA AAG CTA GAT CAC TGT AAC	939
10	Phe Lys Asn Arg Glu Leu Gln Ile Met Arg Lys Leu Asp His Cys Asn	
	95 100 105	
	ATA GTC CGA TTG CGT TAT TTC TTC TAC TCC AGT GGT GAG AAG AAA GAT	987
15	Ile Val Arg Leu Arg Tyr Phe Phe Tyr Ser Ser Gly Glu Lys Lys Asp	
	110 115 120	
	GAG GTC TAT CTT AAT CTG GTG CTG GAC TAT GTT CCG GAA ACA GTA TAC	1035
20	Glu Val Tyr Leu Asn Leu Val Leu Asp Tyr Val Pro Glu Thr Val Tyr	
	125 130 135 140	
	AGA GTT GCC AGA CAC TAT AGT CGA GCC AAA CAG ACG CTC CCT GTG ATT	1083
25	Arg Val Ala Arg His Tyr Ser Arg Ala Lys Gln Thr Leu Pro Val Ile	
	145 150 155	
	TAT GTC AAG TTG TAT ATG TAT CAG CTG TTC CGA AGT TTA GCC TAT ATC	1131
30	Tyr Val Lys Leu Tyr Met Tyr Gln Leu Phe Arg Ser Leu Ala Tyr Ile	
	160 165 170	
	CAT TCC TTT GGA ATC TGC CAT CGG GAT ATT AAA CCG CAG AAC CTC TTG	1179
35	His Ser Phe Gly Ile Cys His Arg Asp Ile Lys Pro Gln Asn Leu Leu	
	175 180 185	
	TTG GAT CCT GAT ACT GCT GTA TTA AAA CTC TGT GAC TTT GGA AGT GCA	1227
40	Leu Asp Pro Asp Thr Ala Val Leu Lys Leu Cys Asp Phe Gly Ser Ala	
	190 195 200	
	AAG CAG CTG GTC CGA GGA GAA CCC AAT GTT TCG TAT ATC TGT TCT CGG	1275
45	Lys Gln Leu Val Arg Gly Glu Pro Asn Val Ser Tyr Ile Cys Ser Arg	
	205 210 215 220	
	TAC TAT AGG GCA CCA GAG TTG ATC TTT GGA GCC ACT GAT TAT ACC TCT	1323
50	Tyr Tyr Arg Ala Pro Glu Leu Ile Phe Gly Ala Thr Asp Tyr Thr Ser	
	225 230 235	

55

AGT ATA GAT GTA TGG TCT GCT GGC TGT GTG TTG GCT GAG CTG TTA CTA 1371  
 Ser Ile Asp Val Trp Ser Ala Gly Cys Val Leu Ala Glu Leu Leu Leu  
 5                   240                   245                   250  
 GGA CAA CCA ATA TTT CCA GGG GAT AGT GGT GTG GAT CAG TTG GTA GAA 1419  
 Gly Gln Pro Ile Phe Pro Gly Asp Ser Gly Val Asp Gln Leu Val Glu  
 10                   255                   260                   265  
 ATA ATC AAG GTC CTG GGA ACT CCA ACA AGG GAG CAA ATC AGA GAA ATG 1467  
 Ile Ile Lys Val Leu Gly Thr Pro Thr Arg Glu Gln Ile Arg Glu Met  
 15                   270                   275                   280  
 AAC CCA AAC TAC ACA GAA TTT AAA TTC CCT CAA ATT AAG GCA CAT CCT 1515  
 Asn Pro Asn Tyr Thr Glu Phe Lys Phe Pro Gln Ile Lys Ala His Pro  
 20                   285                   290                   295                   300  
 TGG ACT AAG GTC TTC CGA CCC CGA ACT CCA CCG GAG GCA ATT GCA CTG 1563  
 Trp Thr Lys Val Phe Arg Pro Arg Thr Pro Pro Glu Ala Ile Ala Leu  
 25                   305                   310                   315  
 TGT AGC CGT CTG CTG GAG TAT ACA CCA ACT GCC CGA CTA ACA CCA CTG 1611  
 Cys Ser Arg Leu Leu Glu Tyr Thr Pro Thr Ala Arg Leu Thr Pro Leu  
 30                   320                   325                   330  
 GAA GCT TGT GCA CAT TCA TTT TTT GAT GAA TTA CCG GAC CCA AAT GTC 1659  
 Glu Ala Cys Ala His Ser Phe Phe Asp Glu Leu Arg Asp Pro Asn Val  
 35                   335                   340                   345  
 AAA CTA CCA AAT GGG CGA GAC ACA CCT GCA CTC TTC AAC TTC ACC ACT 1707  
 Lys Leu Pro Asn Gly Arg Asp Thr Pro Ala Leu Phe Asn Phe Thr Thr  
 40                   350                   355                   360  
 CAA GAA CTG TCA AGT AAT CCA CCT CTG GCT ACC ATC CTT ATT CCT CCT 1755  
 Gln Glu Leu Ser Ser Asn Pro Pro Leu Ala Thr Ile Leu Ile Pro Pro  
 45                   365                   370                   375                   380  
 50  
 55



CAT GCT CGG ATT CAA GCA GCT GCT TCA ACC CCC ACA AAT GCC ACA GCA 1803  
 His Ala Arg Ile Gln Ala Ala Ala Ser Thr Pro Thr Asn Ala Thr Ala  
 5 385 390 395  
 GCG TCA GAT GCT AAT ACT GGA GAC CGT GGA CAG ACC AAT AAT GCT GCT 1851  
 10 Ala Ser Asp Ala Asn Thr Gly Asp Arg Gly Gln Thr Asn Asn Ala Ala  
 400 405 410  
 TCT GCA TCA GCT TCC AAC TCC ACC TGA ACAGTC CCGAGCAGCC AGCTGCACAG 1904  
 15 Ser Ala Ser Ala Ser Asn Ser Thr Stop  
 415 420  
 20 GAAAAACCAC CAGTTACTTG AGTGCTACTC AGCAACACTG GTCACGTTTG GAAAGAATAT 1964  
 TAAAAAGAGA AAAAAATCCT GTTCATTTTA GTGTTCAATT TTTTATTAT TATTGTTGTT 2024  
 25 CTTATTTAAC CTTGTAAAAT ATCTATAAAT ACAAACCAAT TTCATTGTAT TCTCACTTTG 2084  
 AGGG 2088

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## (2) INFORMATION FOR SEQ ID NO:2:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 1932  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: double  
 (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: cDNA to genomic RNA

## (vi) ORIGINAL SOURCE:

- (A) ORGANISM: rat

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:2:

```

GGCCAAGAGA ACGAAGTCTT TTTTTTTTTT TTCTGCGGG AGAACTTAAT GCTGCATTTA   60
TTATTAACCT AGTACCCTAA CATAAAACAA AAGGAAGAAA AGGATTAAGG AAGGAAAAGG   120
TGAATCGAGA AGAGCCATC ATG TCG GGG CGA CCG AGA ACC ACC TCC TTT GCG   172

      Met Ser Gly Arg Pro Arg Thr Thr Ser Phe Ala
      1              5              10

GAG AGC TGC AAG CCA GTG CAG CAG CCT TCA GCT TTT GGT AGC ATG AAA   220
Glu Ser Cys Lys Pro Val Gln Gln Pro Ser Ala Phe Gly Ser Met Lys
      15              20              25

GTT AGC AGA GAT AAA GAT GGC AGC AAG GTA ACC ACA GTG GTG GCA ACT   268
Val Ser Arg Asp Lys Asp Gly Ser Lys Val Thr Thr Val Val Ala Thr
      30              35              40

CCT GGA CAG GGT CCT GAC AGG CCA CAG GAA GTC AGT TAC ACA GAC ACT   316
Pro Gly Gln Gly Pro Asp Arg Pro Gln Glu Val Ser Tyr Thr Asp Thr
      45              50              55

AAA GTC ATT GGA AAT GGG TCA TTT GGT GTG GTA TAT CAA GCC AAA CTT   364
Lys Val Ile Gly Asn Gly Ser Phe Gly Val Val Tyr Gln Ala Lys Leu
      60              65              70              75

```

TGT GAC TCA GGA GAA CTG GTG GCC ATC AAG AAA GTT CTT CAG GAC AAG 412  
 Cys Asp Ser Gly Glu Leu Val Ala Ile Lys Lys Val Leu Gln Asp Lys  
 5 80 85 90  
 CGA TTT AAG AAC CGA GAG CTC CAG ATC ATG AGA AAG CTA GAT CAC TGT 460  
 Arg Phe Lys Asn Arg Glu Leu Gln Ile Met Arg Lys Leu Asp His Cys  
 10 95 100 105  
 AAC ATA GTC CGA TTG CGG TAT TTC TTC TAC TCG AGT GGC GAG AAG AAA 508  
 Asn Ile Val Arg Leu Arg Tyr Phe Phe Tyr Ser Ser Gly Glu Lys Lys  
 15 110 115 120  
 GAT GAG GTC TAC CTT AAC CTG GTG CTG GAC TAT GTT CCG GAA ACA GTG 556  
 Asp Glu Val Tyr Leu Asn Leu Val Leu Asp Tyr Val Pro Glu Thr Val  
 20 125 130 135  
 TAC AGA GTC GCC AGA CAC TAT AGT CGA GCC AAG CAG ACA CTC CCT GTG 604  
 Tyr Arg Val Ala Arg His Tyr Ser Arg Ala Lys Gln Thr Leu Pro Val  
 25 140 145 150 155  
 ATC TAT GTC AAG TTG TAT ATG TAC CAG CTG TTC AGA AGT CTA GCC TAT 652  
 Ile Tyr Val Lys Leu Tyr Met Tyr Gln Leu Phe Arg Ser Leu Ala Tyr  
 30 160 165 170  
 ATC CAT TCC TTT GGG ATC TGC CAT CGA GAC ATT AAA CCA CAG AAC CTC 700  
 Ile His Ser Phe Gly Ile Cys His Arg Asp Ile Lys Pro Gln Asn Leu  
 35 175 180 185  
 TTG CTG GAT CCT GAT ACA GCT GTA TTA AAA CTC TGC GAC TTT GGA AGT 748  
 Leu Leu Asp Pro Asp Thr Ala Val Leu Lys Leu Cys Asp Phe Gly Ser  
 40 190 195 200  
 GCA AAG CAG CTG GTC CGA GGA GAG CCC AAT GTT TCA TAT ATC TGT TCT 796  
 Ala Lys Gln Leu Val Arg Gly Glu Pro Asn Val Ser Tyr Ile Cys Ser  
 45 205 210 215  
 50  
 55

5 CCG TAC TAC AGG GCA CCA GAG CTG ATC TTT GGA GCC ACC GAT TAC ACG 844  
 Arg Tyr Tyr Arg Ala Pro Glu Leu Ile Phe Gly Ala Thr Asp Tyr Thr  
 220 225 230 235  
 10 TCT AGT ATA GAT GTA TGG TCT GCA GGC TGT GTG TTG GCT GAA TTG TTG 892  
 Ser Ser Ile Asp Val Trp Ser Ala Gly Cys Val Leu Ala Glu Leu Leu  
 240 245 250  
 15 CTA GGA CAA CCA ATA TTT CCT GGG GAC AGT GGT GTG GAT CAG TTG GTG 940  
 Leu Gly Gln Pro Ile Phe Pro Gly Asp Ser Gly Val Asp Gln Leu Val  
 255 260 265  
 20 GAA ATA ATA AAG GTC CTA GGA ACA CCA ACA AGG GAG CAA ATT AGA GAA 988  
 Glu Ile Ile Lys Val Leu Gly Thr Pro Thr Arg Glu Gln Ile Arg Glu  
 270 275 280  
 25 ATG AAC CCA AAT TAT ACA GAA TTC AAA TTC CCC CAA ATC AAG GCA CAT 1036  
 Met Asn Pro Asn Tyr Thr Glu Phe Lys Phe Pro Gln Ile Lys Ala His  
 285 290 295  
 30 CCT TGG ACG AAG GTC TTT CGG CCC CGA ACT CCA CCA GAG GCA ATC GCA 1084  
 Pro Trp Thr Lys Val Phe Arg Pro Arg Thr Pro Pro Glu Ala Ile Ala  
 300 305 310 315  
 35 CTG TGT AGC CGT CTC CTG GAG TAC ACG CCG ACC GCC CGG CTA ACA CCA 1132  
 Leu Cys Ser Arg Leu Leu Glu Tyr Thr Pro Thr Ala Arg Leu Thr Pro  
 320 325 330  
 40  
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N : Not identified

## (2) INFORMATION FOR SEQ ID NO:3:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 43  
 (B) TYPE: amino acid  
 (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: peptide

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:3:

Asp Ala Glu Phe Arg His Asp Ser Gly Tyr Glu Val His His Gln

1 5 10 15

Lys Leu Val Phe Phe Ala Glu Asp Val Gly Ser Asn Lys Gly Ala

20 25 30

Ile Ile Gly Leu Met Val Gly Gly Val Val Ile Ala Thr

35 40

## (2) INFORMATION FOR SEQ ID NO:4:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 18  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: other nucleic acid - synthetic DNA  
(sense primer)

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:4:

ATGTCGGGGC GACCGAGA

18

## (2) INFORMATION FOR SEQ ID NO:5:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 18  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: other nucleic acid - synthetic DNA

(antisense primer)

(iv) ANTI-SENCE: yes

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:5:

TCTCGGTCGC CCCGACAT

18

(2) INFORMATION FOR SEQ ID NO:6:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 18
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: other nucleic acid - synthetic DNA  
(sense primer)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:6:

ATGTCAGGGC GGCCCAGA

18

(2) INFORMATION FOR SEQ ID NO:7:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 18
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: other nucleic acid - synthetic DNA  
(anti sense primer)

(iv) ANTI-SENCE: yes

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:7:

TCTGGGCCGC CCTGACAT

18

(2) INFORMATION FOR SEQ ID NO:8:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 170  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: other nucleic acid

(vi) ORIGINAL SOURCE:  
 (A) ORGANISM: rat

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:8:

AAGCTTGTGC ACATTCATTT TTGATGAAT TACGGGACCC AAATGTCAAA CTACCAAATG 60  
 GGCGAGACAC ACCTGCCCTC TTCAACTTTA CCACTCAAGA ACTGTCAAGT AACCCACCTC 120  
 TGGCCACCAT CCTTATCCCT CCTCAGCTC GGATTGAGG AGCTGCTTCA 170

(2) INFORMATION FOR SEQ ID NO:9:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 36  
 (B) TYPE: amino acid  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(vi) ORIGINAL SOURCE:  
 (A) ORGANISM: rat

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:9:

Met Ser Gly Arg Pro Arg Thr Thr Ser Phe Ala Glu Ser Cys Lys Pro  
 1 5 10 15  
 Val Gln Gln Pro Ser Ala Phe Gly Ser Met Lys Val Ser Arg Asp Lys  
 20 25 30  
 Asp Gly Ser Lys  
 35



(2) INFORMATION FOR SEQ ID NO:10:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 34
- (B) TYPE: amino acid
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:10:

Ser Gly Asp Arg Ser Gly Tyr Ser Ser Pro Gly Ser Pro Gly Thr Pro

1 5 10 15

Gly Ser Arg Ser Arg Thr Pro Ser Leu Pro Thr Pro Pro Thr Arg Glu

20 25 30

Pro Lys

(2) INFORMATION FOR SEQ ID NO:11:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 32
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: other nucleic acid - synthetic DNA  
(sense primer)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:11:

TTCGCGAAGA GAGTG C ATATGTCAGG GCGGCC

32

(2) INFORMATION FOR SEQ ID NO:12:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 30
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: other nucleic acid - synthetic DNA  
(antisense primer)

(iv) ANTI-SENCE: yes

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:12:

GGGAGCGTCT GTTGGCTCG ACTATAGTGT

30

(2) INFORMATION FOR SEQ ID NO:13:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 479

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA to genomic RNA

(vi) ORIGINAL SOURCE:

(A) ORGANISM: human being

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:13:

TTCGCGAAGA GAGTGCA TATGTCAGGG CGGCCAGAA CCACCTCCTT TGCGGAGAGC 57  
 TGCAAGCCGG TGCAGCAGCC TTCAGCTTTT GGCAGCATGA AAGTTAGCAG AGACAAGGAC 117  
 GGCAGCAAGG TGACAACAGT GGTGGCAACT CCTGGGCAGG GTCCAGACAG GCCACAAGAA 177  
 GTCAGCTATA CAGACACTAA ACTCATTGGA AATGGATCAT TTGGTGTGGT ATATCAAGCC 237  
 AAACITTTGTG ATTCAGGAGA ACTGGTCGCC ATCAAGAAAG TATTGCAGGA CAAGAGATTT 297  
 AAGAATCGAG AGCTCCAGAT CATGAGAAAG CTAGATCACT GTAACATAGT CCGATTGCGT 357  
 TATTTCTTCT ACTCCAGTGG TGAGAAGAAA GATGAGGTCT ATCTTAATCT GGTGCTGGAC 417  
 TATGTTCCGG AAACAGTATA CAGAGTTGCC AGACACTATA GTCGAGCCAA ACAGACGCTC 477  
 CC 479

#### Claims

1. A preventive or therapeutic agent for Alzheimer's disease which comprises a substance exhibiting an inhibitory action to tau-protein kinase I as an effective component.
2. The preventive or therapeutic agent for Alzheimer's disease according to claim 1 in which the substance exhibiting an inhibitory action to tau-protein kinase I is the one which, when said substance is incubated together with neurons and amymoid beta-protein, has an action of inhibiting the death of said neurons.
3. A preventive or therapeutic agent for Alzheimer's disease which comprises antisense oligonucleotide capable of hybridizing with mRNA or DNA of tau-protein kinase I.
4. The preventive or therapeutic agent for Alzheimer's disease according to claim 3 in which the primary structure of the tau-protein kinase I is expressed by the amino acid sequence which is described in the Seq. ID No. 1 or No. 2 of the attached Sequence Listing.
5. A pharmaceutical composition for prevention or therapy of Alzheimer's disease, which comprises a substance exhibiting an inhibitory action of tau-protein kinase and a pharmaceutically-acceptable

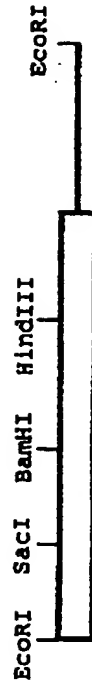
carrier.

6. The pharmaceutical composition according to claim 5 in which the substance exhibiting an inhibitory action to tau-protein kinase I is the one which, when said substance is incubated together with neurons and amyloid beta-protein, has an action of inhibiting the death of said neurons.
7. A pharmaceutical composition for prevention or therapy of Alzheimer's disease, which comprises antisense oligonucleotide capable of hybridizing with mRNA or DNA of tau-protein kinase I.
8. The pharmaceutical composition according to claim 7 in which the primary structure of tau-protein kinase I is represented by the amino acid sequence which is described in the Seq. ID No. 1 or 2 of the attached Sequence Listing.
9. A method for screening an agent useful for a prevention or therapy of Alzheimer's disease, characterized in that amyloid beta-protein, nerve cells and an agent presumed to be effective as a preventive or therapeutic agent for Alzheimer's disease are incubated and, when death of said nerve cells is inhibited, then said agent is judged to be effective as a preventive or therapeutic agent for Alzheimer's disease.
10. The use of a substance exhibiting an inhibitory action of tau-protein kinase I for the manufacture of a medicament for inhibiting the neuronal cell death in the brain characterized in that said substance is applied to the neurons in the brain.
11. The use of an antisense oligonucleotide capable of hybridizing with mRNA or DNA of tau-protein kinase I for the manufacture of a medicament for inhibiting the neuronal cell death in the brain characterized in that said antisense oligonucleotide is applied to the neurons in the brain.
12. The method according to claim 11 in which the primary structure of tau-protein kinase I is expressed by the amino acid sequence described in the Seq. ID No. 1 or No. 2 of the attached Sequence Listing.
13. Tau-Protein kinase I originated from human being, which is characterized by being represented by the amino acid sequence described in the Seq. ID No. 1 of the attached Sequence Listing or the partial sequence thereof.
14. Gene which encodes the human-originated tau-protein kinase I described in claim 13.
15. The gene according to claim 14 in which said gene is represented by the base sequence described in the Seq. ID No. 1 of the attached Sequence Listing.
16. Recombinant human-originated tau-protein kinase I.
17. Recombinant vector which is capable of expressing recombinant human-originated tau-protein kinase I.
18. Transformant which is obtained by subjecting the host cells to a transformation using the recombinant vector of claim 17.
19. A method of producing recombinant human-originated tau-protein kinase I, characterized in that, the transformant of claim 18 is incubated and the recombinant human-originated protein kinase I is collected from its culture.

Fig. 1

Human TPKI cDNA

Clone #1



Clone #2

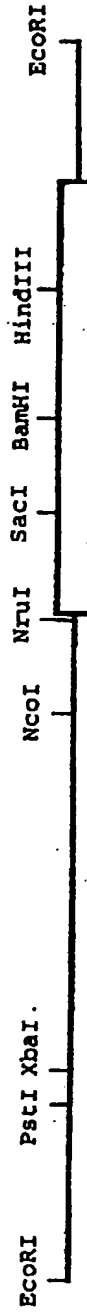


Fig. 2

Comparison of amino acid sequence between human and rat TPKI's

Human	MSGRPRTTSFAESCKPVQPSAFGSMKYSRDKDGSKVTWVATPGQGPDRPQEVSYTDTKLIGNSGFGVV	70
Rat	MSGRPRTTSFAESCKPVQPSAFGSMKYSRDKDGSKVTWVATPGQGPDRPQEVSYTDTKVIGNSGFGVV	70
Human	YQAKLCDSGELVAIKKVLQDKRFKNRELQIMRKLDHCNIVRLRYFFYSYSSGEKKDEVYLNVLDDYVPETVY	140
Rat	YQAKLCDSGELVAIKKVLQDKRFKNRELQIMRKLDHCNIVRLRYFFYSYSSGEKKDEVYLNVLDDYVPETVY	140
Human	RVARHYSRAKQTLPIYVVKLYMYQLFRSLAYIHSFGICHRIKPNQLLLDPDTAVLKLCDFGSAKQLVRG	210
Rat	RVARHYSRAKQTLPIYVVKLYMYQLFRSLAYIHSFGICHRIKPNQLLLDPDTAVLKLCDFGSAKQLVRG	210
Human	EPNVSYSRYYRAPELIFGATDYTSSIDVWSAGCVLAELLGQPIFPDGSVDQLVEIIKVLGTPTREQ	280
Rat	EPNVSYSRYYRAPELIFGATDYTSSIDVWSAGCVLAELLGQPIFPDGSVDQLVEIIKVLGTPTREQ	280
Human	IREMNPNYTEFKFPQIKAHPTWKVFRPRTPEAIALCSRLLLEYTPTARLTPLEACAHSEFFDEL RDPNVKL	350
Rat	IREMNPNYTEFKFPQIKAHPTWKVFRPRTPEAIALCSRLLLEYTPTARLTPLEACAHSEFFDEL RDPNVKL	350
Human	PNGRDTPALFNFTTQELSSNPPLATILIPPHARIQAAASTPTNATAASDANTGDRGQTNNAAASASNSNT	420
Rat	PNGRDTPALFNFTTQELSSNPPLATILIPPHARIQAAASPPANATAASDNTAGDRGQTNNAAASASNSNT	420